WEST Search History

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DB=PGPB,USPT,EPAB; PLUR=YES; OP=ADJ									
	L43	L42 not @py>2000							
	L42	L41 and 15							
. 🔲	L41	L40 and 119	144						
	L40	L39 or l38	11348						
	L39	(514/1,2,423)![CCLS]	10290						
	L38	(424/134.1,144.1)![CCLS]	1142						
	L37	L36 and human	1						
	L36	5444069.pn.	1						
	L35	L34 and 15	5						
	L34	L33 and 125	5						
	L33	(suthanthiran or maluccio).in.	18						
	L32	L31 and losartan	1						
	L31	L30 and 15	1						
	L30	5824696.pn.	1						
	L29	L28 and L21	14						
	L28	L27 not @py>=2000	15						
	L27	L26 and L5	88						
	L26	L25.ab.	1053						
	L25	angiotensin II	7328						
	L24	L22 not @py>2000	80						
	L23	L22 not @ay>2000	119						
	L22	L21 and L20	272						
	L21	antagoni\$ or inhibit\$	665630						
	L20	L18 and L19	286						
	L19	L16.ab.	1946						
	L18	L5 or L6	232985						
	L17	L16 and L5	7934						
	L16	angiotensin\$	15420						
	L15	L6 and L13	0						
· 🗖	L14	L5 and L13	0						

T 12	T 10 T 11 - T 10	
L13	L12 or L11 or L10	31
L12	(5034512 or 4894437 or 5098924 or 5055466 or 4885292 or 5075451 or 4980283 or 5066643).pn.	8
L11	(5114937 or 5106835 or 5063208 or 4845079 or 4845079 or 5089471 or 5071837 or 5064965 or 5063207 or 5036054 or 5036053).pn.	10
L10	(4168267 or 4337201 or 5256687 or 4316906 or 5589499 or 4452740 or 4432970 or 5116835 or 5095119 or 5104869).pn.	13
L9	L4 and L6	1
L8	L5 and L4	1
L7	L5 an dl4	0
L6	angiogen\$ or neovascu\$ or prolifer\$	119958
L5	cancer\$ or tumor\$ or neoplas\$	187735
L4	L3 or L2 or L1	21
L3	4508729.pn.	1
L2	(4316906 or 4374829 or 4344949 or 450879 or 4587256 or 5045553 or 4410520 or 4512924).pn.	9
L1	(5608075 or 5087643 or 6004989 or 6001881 or 598884 or 5234581 or 5889020 or 4473575 or 4105776),pn.	11

END OF SEARCH HISTORY

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     1
                 "Ask CAS" for self-help around the clock
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     2
                New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
NEWS
        DEC 23
                USPAT2
NEWS
        JAN 13
                IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS 5
        JAN 13
                New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
                 INPADOC
        JAN 17
                 Pre-1988 INPI data added to MARPAT
NEWS 6
        JAN 17
                IPC 8 in the WPI family of databases including WPIFV
NEWS
     7
NEWS 8
        JAN 30
                Saved answer limit increased
                 STN AnaVist, Version 1.1, lets you share your STN AnaVist
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        FEB 21
                 visualization results
NEWS 10
        FEB 22
                The IPC thesaurus added to additional patent databases on STN
                Updates in EPFULL; IPC 8 enhancements added
NEWS 11
        FEB 22
NEWS 12
                New STN AnaVist pricing effective March 1, 2006
        FEB 27
NEWS 13
        FEB 28
                MEDLINE/LMEDLINE reload improves functionality
NEWS 14
        FEB 28
                 TOXCENTER reloaded with enhancements
NEWS 15
        FEB 28
                REGISTRY/ZREGISTRY enhanced with more experimental spectral
                 property data
NEWS 16 MAR 01
                 INSPEC reloaded and enhanced
NEWS 17
        MAR 03
                Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 18 MAR 08
                X.25 communication option no longer available after June 2006
NEWS 19 MAR 22
                 EMBASE is now updated on a daily basis
NEWS 20
        APR 03
                New IPC 8 fields and IPC thesaurus added to PATDPAFULL
NEWS 21
        APR 03
                 Bibliographic data updates resume; new IPC 8 fields and IPC
                 thesaurus added in PCTFULL
NEWS 22
        APR 04
                 STN AnaVist $500 visualization usage credit offered
NEWS 23
        APR 12
                 LINSPEC, learning database for INSPEC, reloaded and enhanced
NEWS 24
                 Improved structure highlighting in FQHIT and QHIT display
        APR 12
                 in MARPAT
NEWS 25
        APR 12
                 Derwent World Patents Index to be reloaded and enhanced during
                 second quarter; strategies may be affected
             FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
NEWS EXPRESS
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
              V8.0 AND V8.01 USERS CAN OBTAIN THE UPGRADE TO V8.01a AT
              http://download.cas.org/express/v8.0-Discover/
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http://download.cas.org/express/v8.0-Discover/
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=> file reg COST IN U.S. DOLLARS

SINCE FILE TOTAL
ENTRY SESSION
0.21 0.21

FULL ESTIMATED COST

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TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

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http://www.cas.org/ONLINE/UG/regprops.html

```
=> E "LOSARTAN"/CN 25
                   LOSAN/CN
E1
             1
                   LOSANTIN/CN
E2
             1
             1 --> LOSARTAN/CN
E3
E4
                   LOSARTAN MONOPOTASSIUM SALT/CN
             1
                   LOSARTAN P-TOLUENESULFONATE/CN
E5
             1
                  LOSARTAN POTASSIUM/CN
E6
             1
E7
             1
                   LOSARTAN-HYDROCHLOROTHIAZIDE MIXT./CN
E8
             1
                   LOSBANINE/CN
E9
             1
                   LOSE-URONATE KETOL-ISOMERASE (YERSINIA PESTIS STRAIN CO92 GENE
KDUI)/CN
             1
E10
                   LOSEC/CN
```

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. 1
                   LOSEC SODIUM/CN
 E11
 E12
             1
                   LOSEYITE/CN
 E13
             1
                   LOSFERRON/CN
 E14
             1
                   LOSIGAMONE/CN
 E15
             1
                   LOSIL 1000-50/CN
                   LOSIL 1000-65/CN
 E16
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             1
                  LOSIL 800-50/CN
 E17
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                   LOSINDOLE/CN
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                   LOSK/CN
 E21
             1
                   LOSMIPROFEN/CN
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             1
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 E23
             1
                   LOSOL BLUE/CN
                   LOSOXANTRONE/CN
 E24
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 E25
             1
                   LOSOXANTRONE HYDROCHLORIDE/CN
=> S E3
              1 LOSARTAN/CN
 L1
 => DIS L1 1 SOIDE
 THE ESTIMATED COST FOR THIS REQUEST IS 6.36 U.S. DOLLARS
 DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y) /N:Y
     ANSWER 1 OF 1 REGISTRY COPYRIGHT 2006 ACS on STN
     114798-26-4 REGISTRY
RN
 CN
     1H-Imidazole-5-methanol, 2-butyl-4-chloro-1-[[2'-(1H-tetrazol-5-yl)[1,1'-
     biphenyl]-4-yl]methyl]- (9CI) (CA INDEX NAME)
 OTHER NAMES:
     DUP 89
 CN
 CN
     Lortaan
 CN
      Losartan
 FS
      3D CONCORD
MF
     C22 H23 C1 N6 O
 CI
     COM
 SR
     CA
 LC
      STN Files:
                  ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOSIS,
        BIOTECHNO, CA, CAPLUS, CASREACT, CBNB, CHEMCATS, CHEMINFORMRX, CIN,
        CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*, IMSDRUGNEWS, IMSPATENTS,
        IMSRESEARCH, MEDLINE, MRCK*, PATDPASPC, PROMT, PROUSDDR, RTECS*,
        SCISEARCH, SYNTHLINE, TOXCENTER, USAN, USPATFULL
          (*File contains numerically searchable property data)
      Other Sources:
                      WHO
 DT.CA CAplus document type: Book; Conference; Dissertation; Journal; Patent
        Roles from patents: BIOL (Biological study); PREP (Preparation); PROC
        (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)
       Roles for non-specific derivatives from patents: BIOL (Biological
        study); PREP (Preparation); PRP (Properties); RACT (Reactant or
        reagent); USES (Uses)
       Roles from non-patents: ANST (Analytical study); BIOL (Biological
 RL.NP
        study); FORM (Formation, nonpreparative); MSC (Miscellaneous); OCCU
        (Occurrence); PREP (Preparation); PROC (Process); PRP (Properties); RACT
        (Reactant or reagent); USES (Uses)
 RLD.NP Roles for non-specific derivatives from non-patents: BIOL (Biological
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study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

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2404 REFERENCES IN FILE CA (1907 TO DATE)
31 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
2414 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
=> E "ENALAPRIL"/CN 25
E1
             1
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E2
             1
                   ENALAPRAT/CN
E3
             1 --> ENALAPRIL/CN
                   ENALAPRIL ACETATE/CN
E4
                   ENALAPRIL ACID/CN
E5
             1
             1
                   ENALAPRIL BENZYL ESTER/CN
E6
                   ENALAPRIL CALCIUM DIHYDRATE/CN
E.7
             1
                   ENALAPRIL DIACID/CN
E8
             1
                   ENALAPRIL DIKETOPIPERAZINE/CN
             1
E9
                   ENALAPRIL DKP/CN
E10
             1
                   ENALAPRIL HYDROCHLORIDE/CN
E11
             1
                   ENALAPRIL MALEATE/CN
E12
             1
E13
             1
                   ENALAPRIL MALEATE-HYDROCHLOROTHIAZIDE MIXT./CN
E14
             1
                   ENALAPRIL SODIUM/CN
E15
             1
                   ENALAPRIL TERT-BUTYL ESTER/CN
E16
             1
                   ENALAPRIL-KETANSERIN MIXT./CN
E17
             1
                   ENALAPRILAT/CN
E18
             1
                   ENALAPRILAT SRS/CN
                   ENALAPRILAT-KETANSERIN MIXT./CN
E19
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E20
             1
                   ENALAPRILIC ACID/CN
E21
             1
                   ENALIN A/CN
E22
             1
E23
             1
                   ENALIN B/CN
E24
             1
                    ENALITE/CN
E25
                    ENALKIREN/CN
=> S E3
             1 ENALAPRIL/CN
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=> DIS L2 1 RN

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2006 ACS on STN RN 75847-73-3 REGISTRY

=> file caplus COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 12.51 12.72

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=> s 11 or 12

2414 L1

2695 L2

L3 4684 L1 OR L2

=> s cancer? or tumor? or neoplas?

290705 CANCER?

425403 TUMOR?

446494 NEOPLAS?

L4 704495 CANCER? OR TUMOR? OR NEOPLAS?

=> s 14 and 13

L5 146 L4 AND L3

=> s 13 (L) 14

L6 22 L3 (L) L4

=> s 16 not py>2000

5711376 PY>2000

L7 3 L6 NOT PY>2000

=> d ibib 1-3

L7 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1999:794097 CAPLUS

DOCUMENT NUMBER:

132:18639

TITLE:

Do ACE-inhibitors suppress tumor necrosis factor- α production in advanced chronic renal

failure?

AUTHOR(S):

Stenvinkel, P.; Andersson, P.; Wang, T.; Lindholm, B.;

Bergstrom, J.; Palmblad, J.; Heimburger, O.;

Cederholm, T.

CORPORATE SOURCE:

Departments of Clinical Science, Divisions of Renal

Medicine and Baxter Novum, Stockholm, Swed.

SOURCE:

Journal of Internal Medicine (1999), 246(5), 503-507

CODEN: JINMEO; ISSN: 0954-6820

PUBLISHER:

Blackwell Science Ltd.

DOCUMENT TYPE: LANGUAGE:

Journal English

REFERENCE COUNT:

THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 1999:688629 CAPLUS DOCUMENT NUMBER: 132:175414 TITLE: Pharmacological modifications of the partial pressure of oxygen in murine tumors: evaluation using in vivo EPR oximetry AUTHOR(S): Gallez, Bernard; Jordan, Benedicte F.; Baudelet, Christine; Misson, Pierre-Damien CORPORATE SOURCE: Laboratory of Medicinal Chemistry and Radiopharmacy, Universite Catholique de Louvain, Brussels, Belg. SOURCE: Magnetic Resonance in Medicine (1999), 42(4), 627-630 CODEN: MRMEEN; ISSN: 0740-3194 PUBLISHER: Wiley-Liss, Inc. DOCUMENT TYPE: Journal LANGUAGE: English REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN 1995:845231 CAPLUS ACCESSION NUMBER: 123:275949 DOCUMENT NUMBER: TITLE: Angiotensin-converting-enzyme inhibitors suppress synthesis of tumor necrosis factor and interleukin 1 by human peripheral blood mononuclear cells AUTHOR(S): Schindler, Ralf; Dinarello, Charles A.; Koch, Karl-M. CORPORATE SOURCE: Department of Nephrology, Medical School Hannover, Berlin, D-14050, Germany Cytokine (1995), 7(6), 526-33 SOURCE: CODEN: CYTIE9; ISSN: 1043-4666 PUBLISHER: Academic DOCUMENT TYPE: Journal English LANGUAGE: => d ibibi abs 2 'IBIBI' IS NOT A VALID FORMAT FOR FILE 'CAPLUS' The following are valid formats: ABS ----- GI and AB ALL ----- BIB, AB, IND, RE APPS ----- AI, PRAI BIB ----- AN, plus Bibliographic Data and PI table (default) CAN ----- List of CA abstract numbers without answer numbers CBIB ----- AN, plus Compressed Bibliographic Data CLASS ----- IPC, NCL, ECLA, FTERM DALL, ----- ALL, delimited (end of each field identified) DMAX ----- MAX, delimited for post-processing FAM ----- AN, PI and PRAI in table, plus Patent Family data FBIB ----- AN, BIB, plus Patent FAM IND ---- Indexing data IPC ----- International Patent Classifications MAX ----- ALL, plus Patent FAM, RE PATS ---- PI, SO SAM ----- CC, SX, TI, ST, IT SCAN ----- CC, SX, TI, ST, IT (random display, no answer numbers; SCAN must be entered on the same line as the DISPLAY, e.g., D SCAN or DISPLAY SCAN) STD ----- BIB, CLASS IABS ----- ABS, indented with text labels IALL ----- ALL, indented with text labels IBIB ----- BIB, indented with text labels

IMAX ----- MAX, indented with text labels ISTD ----- STD, indented with text labels

OBIB ----- AN, plus Bibliographic Data (original)

OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations SIBIB ----- IBIB, no citations

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HITIND ----- IC, ICA, ICI, NCL, CC and index field (ST and IT)

containing hit terms

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HITSTR ----- HIT RN, its text modification, its CA index name, and

its structure diagram

HITSEQ ----- HIT RN, its text modification, its CA index name, its

structure diagram, plus NTE and SEQ fields

FHITSTR ---- First HIT RN, its text modification, its CA index name, and

its structure diagram

FHITSEQ ---- First HIT RN, its text modification, its CA index name, its

structure diagram, plus NTE and SEQ fields

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OCC ----- Number of occurrence of hit term and field in which it occurs

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ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN 1.7

ACCESSION NUMBER: 1999:688629 CAPLUS

DOCUMENT NUMBER:

132:175414

TITLE:

Pharmacological modifications of the partial pressure

of oxygen in murine tumors: evaluation using in vivo

EPR oximetry

Gallez, Bernard; Jordan, Benedicte F.; Baudelet, AUTHOR(S):

Christine; Misson, Pierre-Damien

CORPORATE SOURCE: Laboratory of Medicinal Chemistry and Radiopharmacy,

Universite Catholique de Louvain, Brussels, Belg.

Magnetic Resonance in Medicine (1999), 42(4), 627-630

CODEN: MRMEEN; ISSN: 0740-3194

PUBLISHER: Wiley-Liss, Inc.

DOCUMENT TYPE: Journal English LANGUAGE:

REFERENCE COUNT: THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS 12

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d ibib abs 2

SOURCE:

ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1999:688629 CAPLUS

DOCUMENT NUMBER: 132:175414

TITLE: Pharmacological modifications of the partial pressure

of oxygen in murine tumors: evaluation using in vivo

EPR oximetry

AUTHOR(S): Gallez, Bernard; Jordan, Benedicte F.; Baudelet,

Christine; Misson, Pierre-Damien

CORPORATE SOURCE: Laboratory of Medicinal Chemistry and Radiopharmacy,

Universite Catholique de Louvain, Brussels, Belg.

SOURCE: Magnetic Resonance in Medicine (1999), 42(4), 627-630

CODEN: MRMEEN; ISSN: 0740-3194

PUBLISHER: Wiley-Liss, Inc.

DOCUMENT TYPE: Journal LANGUAGE: English

AB EPR oximetry using an implantable paramagnetic probe was used to quantify the partial pressure of O (pO2) in tissues in a transplantable mouse tumor model after administration of 34 different vasodilators belonging to one of the following classes: angiotensin-converting enzyme inhibitors, Ca2+ antagonists, α -antagonists, K+ channel openers, β -blockers, NO donors, and peripheral vasoactive agents. Twenty-four compds. were efficient in significantly increasing the local pO2 in a majority of tumors. The increase of local pO2 by the pharmacol. treatments was lower than that achieved by using O or carbogen breathing. This technique offers a tool for rapidly and accurately measuring treatment-induced modifications of pO2 in tumors.

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s prolif? or angiogen? or neovascular?

240616 PROLIF?

35334 ANGIOGEN?

7034 NEOVASCULAR?

L8 266785 PROLIF? OR ANGIOGEN? OR NEOVASCULAR?

=> d his

(FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)

FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006

E "LOSARTAN"/CN 25

L1 1 S E3

E "ENALAPRIL"/CN 25

L2 1 S E3

FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006

L3 4684 S L1 OR L2

L4 704495 S CANCER? OR TUMOR? OR NEOPLAS?

L5 146 S L4 AND L3

L6 22 S L3 (L) L4

L7 3 S L6 NOT PY>2000

L8 266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?

=> s 18 and 13

L9 220 L8 AND L3

=> s 18 (1) 13

L10 46 L8 (L) L3

=> s 110 not py>2000

5711376 PY>2000

L11 18 L10 NOT PY>2000

=> d ibib 1-6

L11 ANSWER 1 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:23055 CAPLUS

DOCUMENT NUMBER: 134:161364

TITLE: Retinal neovascularization is prevented by blockade of

the renin-angiotensin system

AUTHOR(S): Moravski, Christina J.; Kelly, Darren J.; Cooper, Mark

E.; Gilbert, Richard E.; Bertram, John F.; Shahinfar,

Shahnaz; Skinner, Sandford L.; Wilkinson-Berka,

Jennifer L.

CORPORATE SOURCE: Department of Physiology, The University of Melbourne,

Parkville, 3010, Australia

Hypertension (2000), 36(6), 1099-1104 SOURCE:

CODEN: HPRTDN; ISSN: 0194-911X Lippincott Williams & Wilkins

DOCUMENT TYPE: Journal

PUBLISHER:

LANGUAGE: English

REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 2 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

2000:871260 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 135:322

TITLE: The effect of losartan on intimal thickening in rats

after vascular balloon injury

AUTHOR(S): Li, Yonghong; Dong, Guoxiong; Guo, Minglei

Department of Emergency, The Affiliated Hospital of CORPORATE SOURCE:

Qingdao University Medical College, Tsingtao, 266003,

Peop. Rep. China

SOURCE: Qingdao Daxue Yixueyuan Xuebao (2000), 36(4), 271-273

CODEN: ODYXAE

Qingdao Daxue Yixueyuan Xuebao Bianjibu PUBLISHER:

Journal DOCUMENT TYPE: Chinese LANGUAGE:

L11 ANSWER 3 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

2000:808012 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 134:290166

TITLE: Antagonistic effect of monocyte chemotactic protein-1

monoclonal antibody and losartan on proliferation and

migration of Ang-II-mediated VSMCs in vitro

AUTHOR(S): He, Zuo-yun; Li, Ai-min

Department of Cardiology, Xinqiao Hospital, Third CORPORATE SOURCE:

Military Medical University, Chungking, 400037, Peop.

Rep. China

Di-San Junyi Daxue Xuebao (2000), 22(9), 815-818 SOURCE:

CODEN: DYXUE8; ISSN: 1000-5404

PUBLISHER: Di-San Junyi Daxue

DOCUMENT TYPE: Journal LANGUAGE: Chinese

L11 ANSWER 4 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:413731 CAPLUS

DOCUMENT NUMBER: 133:246971

The stimulating effects of neuropeptide Y on cultured TITLE:

arterial smooth muscle cell proliferation and losartan

treatment

Huang, Shao-Hua; Liu, Jian-Kang; Chen, Min-Sheng AUTHOR(S):

First Affiliated Hospital, Guangzhou Medical College, CORPORATE SOURCE:

Canton, 510182, Peop. Rep. China

SOURCE: Zhongguo Bingli Shengli Zazhi (2000), 16(3), 211-213

CODEN: ZBSZEB; ISSN: 1000-4718

PUBLISHER: Jinan Daxue DOCUMENT TYPE: Journal LANGUAGE: Chinese

L11 ANSWER 5 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:307168 CAPLUS

DOCUMENT NUMBER: 133:187796

TITLE: Enalapril inhibits growth and proliferation of various

tissues in rat normotensive four-sixths kidney

ablation nephropathy

AUTHOR(S): Gajdos, Martin; Krivosikova, Zora; Sebekova, Katarina;

Lajdova, Ingrid; Spustova, Viera; Dzurik, Rastislav

CORPORATE SOURCE: Institute of Preventive and Clinical Medicine,

Bratislava, 833 01, Slovakia

SOURCE: Kidney & Blood Pressure Research (2000), 23(2),

106-112

CODEN: KBPRFC; ISSN: 1420-4096

PUBLISHER: S. Karger AG

DOCUMENT TYPE: Journal LANGUAGE: English

REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 6 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:52976 CAPLUS

DOCUMENT NUMBER: 132:329674

TITLE: Enalapril in subantihypertensive dosage attenuates

kidney proliferation and functional recovery in normotensive ablation nephropathy of the rat

AUTHOR(S): Krivosikova, Z.; Sebekova, K.; Spustova, V.; Lajdova,

I.; Dzurik, R.

CORPORATE SOURCE: Institute of Preventive and Clinical Medicine,

Bratislava, Slovakia

SOURCE: Physiological Research (Prague) (1999), 48(6), 429-435

CODEN: PHRSEJ; ISSN: 0862-8408

PUBLISHER: Institute of Physiology, Academy of Sciences of the

Czech Republic

DOCUMENT TYPE: Journal LANGUAGE: English

REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d ibib 7-12

L11 ANSWER 7 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:14644 CAPLUS

DOCUMENT NUMBER: 132:288508

TITLE: Smooth muscle cell proliferation in the ductus

arteriosus and the descending aorta, and effects of enalapril on SMC proliferation in perinatal rats

AUTHOR(S): Takizawa, Tatsuya; Kawahata, Mariko; Ikeda, Yoshinori;

Yamamoto, Masako; Arishima, Kazuyoshi; Muto, Makoto;

Masaoka, Toshio

CORPORATE SOURCE: Departments of Developmental and Reproductive

Biotechnology, Azabu University School of Veterinary

Medicine, Sagamihara, 229-8501, Japan

SOURCE: Journal of Veterinary Medical Science (1999), 61(11),

1215-1218

CODEN: JVMSEQ; ISSN: 0916-7250

PUBLISHER: Japanese Society of Veterinary Science

DOCUMENT TYPE: Journal LANGUAGE: English

REFERENCE COUNT: 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1999:649604 CAPLUS

DOCUMENT NUMBER: 132:178991

TITLE: Effect of angiotensin II receptor antagonist on

proliferation of glomerular mesangial cells from

spontaneously hypertensive rats

AUTHOR(S): Chen, Shaoxing; Guo, Jizhen; Qiang, Weiguo; Du, Jian;

Liu, Xiaoping; Zhu, Dingliang

CORPORATE SOURCE: Department of Hypertension, Shanghai Institute of

Hypertension, Ruijin Hospital, Shanghai Second Medical

University, Shanghai, 200025, Peop. Rep. China

SOURCE: Shanghai Dier Yike Daxue Xuebao (1999), 19(4), 292-294

CODEN: SDDXE3; ISSN: 0258-5898

PUBLISHER: Shanghai Dier Yike Daxue Xuebao Bianjibu

DOCUMENT TYPE: Journal LANGUAGE: Chinese

L11 ANSWER 9 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1999:360095 CAPLUS

DOCUMENT NUMBER: 131:179549

TITLE: Angiotensin II receptor antagonists prevent neointimal

proliferation in a porcine coronary artery organ

culture model

AUTHOR(S): Wilson, David P.; Saward, Laura; Zahradka, Peter; Kee

Cheung, Po

CORPORATE SOURCE: Department of Physiology, University of Manitoba, MB,

Can.

SOURCE: Cardiovascular Research (1999), 42(3), 761-772

CODEN: CVREAU; ISSN: 0008-6363

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

REFERENCE COUNT: 61 THERE ARE 61 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 10 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1999:306644 CAPLUS

DOCUMENT NUMBER: 131:68521

TITLE: Effects of TH-142177 on angiotensin II-induced

proliferation, migration and intracellular signaling in vascular smooth muscle cells and on neointimal

thickening after balloon injury

AUTHOR(S): Nozawa, Yoshihisa; Matsuura, Naosuke; Miyake,

Hidekazu; Yamada, Shizuo; Kimura, Ryohei

CORPORATE SOURCE: Pharmacology Research Laboratory, Taiho Pharmaceutical

Co., Ltd., Tokushima, 771-0194, Japan Life Sciences (1999), 64(22), 2061-2070

CODEN: LIFSAK; ISSN: 0024-3205

PUBLISHER: Elsevier Science Inc.

DOCUMENT TYPE: Journal LANGUAGE: English

SOURCE:

REFERENCE COUNT: 31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 11 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:284031 CAPLUS

DOCUMENT NUMBER: 129:36278

TITLE: Divergent effects of angiotensin-converting enzyme

inhibition and angiotensin II-receptor antagonism on

myocardial cellular proliferation and collagen deposition after myocardial infarction in rats

AUTHOR(S): Taylor, Kenneth; Patten, Richard D.; Smith, John J.;

Aronovitz, Mark J.; Wight, Joseph; Salomon, Robert N.;

Konstam, Marvin A.

CORPORATE SOURCE: Division of Cardiology, Department of Medicine, New

England Medical Center, Tufts University School of

Medicine, Boston, MA, 02111, USA

SOURCE: Journal of Cardiovascular Pharmacology (1998), 31(5),

654-660

CODEN: JCPCDT; ISSN: 0160-2446 Lippincott-Raven Publishers

PUBLISHER: Lippincott-H DOCUMENT TYPE: Journal

DOCUMENT TYPE: Journal LANGUAGE: English

REFERENCE COUNT: 42 THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 12 OF 18 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1997:108045 CAPLUS

DOCUMENT NUMBER: 126:195036

TITLE: In vitro and in vivo effects of UP 269-6, a new potent

orally active nonpeptide angiotensin II receptor

antagonist, on vascular smooth muscle cell

proliferation

AUTHOR(S): Virone-Oddos, A.; Desangle, V.; Provost, D.; Cazes,

M.; Caussade, F.; Cloarec, A.

CORPORATE SOURCE: Lab. UPSA, Rueil-Malmaison, 92506, Fr.

SOURCE: British Journal of Pharmacology (1997), 120(3),

488-494

CODEN: BJPCBM; ISSN: 0007-1188

PUBLISHER: Stockton
DOCUMENT TYPE: Journal
LANGUAGE: English

=> d his

L3

(FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)

FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006

E "LOSARTAN"/CN 25

L1 1 S E3

E "ENALAPRIL"/CN 25

L2 1 S E3

FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006

4684 S L1 OR L2

L4 704495 S CANCER? OR TUMOR? OR NEOPLAS?

L5 146 S L4 AND L3 L6 22 S L3 (L) L4

L7 3 S L6 NOT PY>2000

L8 266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?

L9 220 S L8 AND L3 L10 46 S L8 (L) L3

L11 18 S L10 NOT PY>2000

=> s 110 and 14

L12 7 L10 AND L4

=> d ibib 1-7

L12 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2006:182557 CAPLUS

DOCUMENT NUMBER: 144:267234

TITLE: Effects of angiotensin II receptor antagonist,

Losartan on the apoptosis, proliferation and migration

of the human pancreatic stellate cells

AUTHOR(S): Liu, Wen-Bin; Wang, Xing-Peng; Wu, Kai; Zhang, Ru-Ling CORPORATE SOURCE: Shanghai No. 1 People's Hospital, Shanghai Jiaotong

University, Shanghai, 200080, Peop. Rep. China SOURCE:

World Journal of Gastroenterology (2005), 11(41),

6489-6494

CODEN: WJGAF2; ISSN: 1007-9327 PUBLISHER: World Journal of Gastroenterology

DOCUMENT TYPE: Journal English LANGUAGE:

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

2005:577447 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 143:379379

Effects of Combined Endothelin and Angiotensin II TITLE:

> Antagonism on Growth Factor-Induced Proliferation of Vascular Smooth Muscle Cells Isolated from Uremic Rats

Wolf, Sabine; Sauter, Gabriele; Risler, Teut; Brehm, AUTHOR(S):

Bernhard

CORPORATE SOURCE: Medical Clinic IV, Department of Hypertension and

Renal Failure and Endocrinology, University of

Tuebingen, Tuebingen, Germany

Renal Failure (2005), 27(4), 465-474 SOURCE:

CODEN: REFAE8; ISSN: 0886-022X

Taylor & Francis, Inc. PUBLISHER:

Journal DOCUMENT TYPE: English LANGUAGE:

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

2005:291659 CAPLUS ACCESSION NUMBER:

143:323408 DOCUMENT NUMBER:

Blockage of angiotensin II type I receptor decreases TITLE:

the synthesis of growth factors and induces apoptosis

in C6 cultured cells and C6 rat glioma Arrieta, O.; Guevara, P.; Escobar, E.;

Garcia-Navarrete, R.; Pineda, B.; Sotelo, J.

Neuroimmunology Unit of the National Institute of CORPORATE SOURCE:

Neurology and Neurosurgery of Mexico, Mexico City,

14269, Mex.

British Journal of Cancer (2005), 92(7), 1247-1252 SOURCE:

CODEN: BJCAAI; ISSN: 0007-0920

Nature Publishing Group PUBLISHER:

Journal DOCUMENT TYPE: LANGUAGE: English

AUTHOR(S):

CORPORATE SOURCE:

THERE ARE 55 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 55

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

2005:88914 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 142:403819

Influence of growth factors on the proliferation of TITLE: vascular smooth muscle cells isolated from subtotally

nephrectomized rats after endothelin or angiotensin II

antagonism

Wolf, Sabine C.; Sauter, Gabriele; Rodemann, AUTHOR(S):

Hans-Peter; Risler, Teut; Brehm, Bernhard R. Medical Clinic III, Department of Cardiology, Nephrology, Hypertension and Renal Failure,

Eberhard-Karls-University, Tuebingen, D-72076, Germany

Nephrology, Dialysis, Transplantation (2005), 20(2), SOURCE:

312-318

CODEN: NDTREA; ISSN: 0931-0509

PUBLISHER: Oxford University Press

DOCUMENT TYPE: Journal LANGUAGE: English

THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 21 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

2003:777526 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 139:286322

TITLE: PAR receptor-mediated antiangiogenic activity of

thrombin and use of PAR receptor agonists for the

treatment of cancer and other angiogenesis-associated diseases

Sukhatme, Vikas P.; Merchan, Jaime; Chan, Barden INVENTOR(S):

PATENT ASSIGNEE(S): Beth Israel Deaconess Medical Center, USA

SOURCE: PCT Int. Appl., 77 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.				KIND DATE			APPLICATION NO.						DATE				
	WO	2003	0799	78		A2		2003	1002	1	WO 2	003-1	US812	21		2	0030	314
•	WO 2003079978				A3 20040226													
		W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
			co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NI,	NO,	NZ,	OM,
			PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,
			TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW					
		RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
			KG,	ΚZ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
•			FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,
			BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG
AU 2003218213				A1		20031008 AU 2003-218213						20030314						
US 2005232925					A1		20051020			US 2005-508317				20050616				
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										1	WO 2	003-1	US81:	21	1	W 2	0030	314

L12 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2003:272945 CAPLUS

DOCUMENT NUMBER: 139:301459

TITLE: The effect of commonly used drugs on angiogenesis Sartippour, Maryam R.; De Leon, Ernesto; Rubio, AUTHOR(S):

Rosalio; Brooks, Mai N.

CORPORATE SOURCE: Department of Surgery, Division of Oncology,

University of California, Los Angeles, CA, 90095, USA

SOURCE: Anticancer Research (2003), 23(1A), 231-234

CODEN: ANTRD4; ISSN: 0250-7005

PUBLISHER: International Institute of Anticancer Research

Journal DOCUMENT TYPE: English LANGUAGE:

THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 17 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:14362 CAPLUS

DOCUMENT NUMBER: 136:379590

AT1 receptor is present in glioma cells; its blockage TITLE:

reduces the growth of rat glioma

AUTHOR(S): Rivera, E.; Arrieta, O.; Guevara, P.; Duarte-Rojo, A.; Sotelo, J.

CORPORATE SOURCE:

Neuroimmunology Unit, National Institute of Neurology

and Neurosurgery of Mexico, Mexico, 14269, Mex. British Journal of Cancer (2001), 85(9), 1396-1399

SOURCE: CODEN: BJCAAI; ISSN: 0007-0920

PUBLISHER:

Harcourt Publishers Ltd.

DOCUMENT TYPE:

Journal English

LANGUAGE: REFERENCE COUNT:

34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s losartan or enalapril

4562 LOSARTAN

1 LOSARTANS

4562 LOSARTAN

(LOSARTAN OR LOSARTANS)

3733 ENALAPRIL

7727 LOSARTAN OR ENALAPRIL L13

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(FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)

FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006

E "LOSARTAN"/CN 25

L1 1 S E3

E "ENALAPRIL"/CN 25

L2 1 S E3

FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006

L3 4684 S L1 OR L2

704495 S CANCER? OR TUMOR? OR NEOPLAS? L4

146 S L4 AND L3 L5

L6 22 S L3 (L) L4

L7 3 S L6 NOT PY>2000

L8 266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?

L9 220 S L8 AND L3

L1046 S L8 (L) L3

18 S L10 NOT PY>2000 L11

L12 7 S L10 AND L4

L13 7727 S LOSARTAN OR ENALAPRIL

=> s 113 (1) 14

92 L13 (L) L4

=> s 114 not py>2000

5711376 PY>2000

28 L14 NOT PY>2000 L15

=> s 114 not py>1999

6603140 PY>1999

23 L14 NOT PY>1999 L16

=> s 116 and 18

4 L16 AND L8

=> d ibib 1-4

L17 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

1998:204801 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 128:304377

TITLE: Angiotensin II-induced Ca2+ mobilization and prolactin

release in normal and hyperplastic pituitary cells AUTHOR(S): Diaz-Torga, Graciela; Gonzelez Iglesias, Arturo;

Achaval-Zaia, Rita; Libertun, Carlos; Becu-Villalobos,

Damasia

CORPORATE SOURCE: Inst. Biologia Medicina Experimental, Consejo

Investigaciones Cientificas Tecnicas, Buenos Aires,

1428, Argent.

SOURCE: American Journal of Physiology (1998), 274(3, Pt. 1),

E534-E540

CODEN: AJPHAP; ISSN: 0002-9513 American Physiological Society

DOCUMENT TYPE:

Journal

PUBLISHER: LANGUAGE:

English

REFERENCE COUNT:

31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L17 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1997:256278 CAPLUS

DOCUMENT NUMBER:

126:329068

TITLE:

Sequential development of angiotensin receptors and

angiotensin I converting enzyme during

angiogenesis in the rat subcutaneous sponge

granuloma

Walsh, David A.; Hu, De-En; Wharton, John; Catravas, AUTHOR(S):

John D.; Blake, David R.; Fan, Tai-Ping D.

Inflammation Group, London Hospital Medical College, CORPORATE SOURCE:

London, E1 2AD, UK

British Journal of Pharmacology (1997), 120(7), SOURCE:

1302-1311

CODEN: BJPCBM; ISSN: 0007-1188

PUBLISHER: DOCUMENT TYPE:

LANGUAGE:

Stockton Journal English

L17 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1995:939339 CAPLUS

DOCUMENT NUMBER:

124:52845

TITLE:

Is estrogen-induced pituitary hyperplasia and hyperprolactinemia mediated by angiotensin II?

AUTHOR(S):

Pawlikowski, M.; Mucha, S.; Kunert-Radek, J.; Sepien,

H.; Pisarek, H.; Stawowy, A.

CORPORATE SOURCE:

Institute Endocrinology, Medical University Lodz,

Lodz, Pol.

SOURCE:

Advances in Experimental Medicine and Biology (1995),

377 (Tissue Renin-Angiotensin Systems), 371-8

CODEN: AEMBAP; ISSN: 0065-2598

PUBLISHER:

Plenum Journal English

DOCUMENT TYPE: LANGUAGE:

L17 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

CORPORATE SOURCE:

1995:845231 CAPLUS 123:275949

DOCUMENT NUMBER: TITLE:

Angiotensin-converting-enzyme inhibitors suppress synthesis of tumor necrosis factor and interleukin 1

by human peripheral blood mononuclear cells

AUTHOR(S):

Schindler, Ralf; Dinarello, Charles A.; Koch, Karl-M. Department of Nephrology, Medical School Hannover,

Berlin, D-14050, Germany

SOURCE:

Cytokine (1995), 7(6), 526-33 CODEN: CYTIE9; ISSN: 1043-4666

PUBLISHER: Academic Journal DOCUMENT TYPE:

LANGUAGE: English

=> d kwic 1-2

- L17 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN . . . on prolactin release, intracellular calcium ([Ca2+]i) AB mobilization, and [3H]thymidine uptake in cells from normal rat pituitaries and from estrogen-induced pituitary tumors. ANG II (10-7 to 10-9 M) increased prolactin release significantly in control and not in tumoral cells. In control cells, ANG II (10-6 to 10-9 M) produced an immediate spike of [Ca2+]i followed by a plateau.. significantly between 10-10 and 10-8 M ANG II, whereas the onset of the spike was retarded with decreasing concns. In tumoral cells, ANG II did not produce a spike phase even at 10-6 M. ANG II-induced prolactin release and calcium mobilization were blocked by losartan (AT1 receptor antagonist) and not by PD-123319 (AT2 antagonist). Finally, [3H]thymidine uptake was not modified by ANG II (10-7 to. . . estrogenic treatment alters in vitro pituitary response to ANG II. Alterations might function to limit excessive prolactin secretion of hypersecreting tumors. Besides, ANG II does not modify DNA synthesis in vitro of cells from normal or tumor -derived hypophyses.
- IT Cell proliferation
 Pituitary gland
 (angiotensin II-induced Ca2+ mobilization and prolactin release in normal and hyperplastic pituitary cells)
- L17 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

 TI Sequential development of angiotensin receptors and angiotensin I converting enzyme during angiogenesis in the rat subcutaneous sponge granuloma
- The vasoconstrictor peptide angiotensin II (AII) can stimulate AB angiogenesis, an important process in wound healing, tumor growth and chronic inflammation. To elucidate mechanisms underlying AII-enhanced angiogenesis, we have studied a s.c. sponge granuloma model in the rat by use of 133Xe clearance, morphometry and quant. in. . . directly into the sponge, AII (1 nmol day-1) increased 133Xe clearance from, and fibrovascular growth in sponge granulomas, indicating enhanced angiogenesis 6 to 12 days after implantation. This AII-enhanced angiogenesis was inhibited by daily doses (100 nmol/sponge) of the specific but subtype non-selective AII receptor antagonist (Sarl, Ile8)AII, and by the selective non-peptide AT1 receptor antagonists losartan and DuP 532. In contrast, AII-enhanced neovascularization was not inhibited by the AT2 receptor antagonist PD123319, nor was it mimicked by the AT2 receptor agonist CGP42112A (each. . . (ACE) inhibitors captopril (up to 100 μg/sponge day-1) and lisinopril (40 μg/sponge day-1), or AII receptor antagonists did not affect angiogenesis in the absence of exogenous AII. [1251]-(Sarl, Ile8)AII binding sites with characteristics of AT1 receptors were localized to microvessels and. days after sponge implantation. [1251]-351A bound less densely to sponge stroma than to skin. We propose that AII can stimulate angiogenesis, acting via AT1 receptors within the sponge granuloma. AT1 and AT2 receptors and ACE develop sequentially during microvascular maturation, and the role of the endogenous angiotensin system in angiogenesis will depend on the balanced local expression of its various components. Pharmacol. modulation of this balance may provide novel therapeutic approaches in angiogenesis -dependent diseases.
- ST angiotensin receptor ACE angiogenesis
- IT Angiotensin receptors
 RL: BAC (Biological activity or effector, except adverse); BPR (Biological

process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(AT1; sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma)

IT Angiotensin receptors

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(AT2; sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma)

IT Disease, animal

(angiogenesis-dependent; sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma in relation to treatment of angiogenesis -dependent diseases)

IT Angiogenesis

(neovascularization; sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma)

IT Granuloma

(sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma)

IT Angiogenesis

(sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma in relation to treatment of angiogenesis-dependent diseases)

IT 9015-82-1, Angiotensin I converting enzyme 11128-99-7, Angiotensin II RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(sequential development of angiotensin receptors and ACE during angiogenesis in rat s.c. sponge granuloma)

=> d kwic 3

L17 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

AB . . . role of angiotensin II (AII) in estrogen action on the rat anterior pituitary gland. AII is able to enhance the proliferation of cells isolated from estrogen-induced pituitary tumors, as well as cells isolated from human pituitary adenomas. Angiotensin-converting enzyme inhibitors enalapril and enalprilate decreased the cell proliferation indexes of DES-induced pituitary tumors. AII receptor blockers diminished the d. of prolactin-immunoreactive cells in DES-induced pituitary

=> d 3 abs

tumors.

L17 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

AB A study was done to investigate the role of angiotensin II (AII) in estrogen action on the rat anterior pituitary gland. AII is able to enhance the proliferation of cells isolated from estrogen-induced pituitary tumors, as well as cells isolated from human pituitary adenomas. Angiotensin-converting enzyme inhibitors enalapril and enalprilate decreased the cell proliferation indexes of DES-induced pituitary tumors. AII receptor blockers diminished the d. of prolactin-immunoreactive cells in DES-induced pituitary tumors.

(FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)

FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006 E "LOSARTAN"/CN 25 1 S E3 L1 E "ENALAPRIL"/CN 25 L2 1 S E3 FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006 4684 S L1 OR L2 L3L4704495 S CANCER? OR TUMOR? OR NEOPLAS? L5 146 S L4 AND L3 L6 22 S L3 (L) L4 L7 3 S L6 NOT PY>2000 266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR? L8 L9 220 S L8 AND L3 46 S L8 (L) L3 L10 18 S L10 NOT PY>2000 L117 S L10 AND L4 L12 7727 S LOSARTAN OR ENALAPRIL L13 92 S L13 (L) L4 28 S L14 NOT PY>2000 L15 23 S L14 NOT PY>1999 L16 L17 4 S L16 AND L8 => file pctfull COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 73.64 86.36 SINCE FILE DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) TOTAL ENTRY SESSION CA SUBSCRIBER PRICE -3.75-3.75FILE 'PCTFULL' ENTERED AT 10:25:28 ON 26 APR 2006 COPYRIGHT (C) 2006 Univentio 25 APR 2006 <20060425/UP> FILE LAST UPDATED: 200616 <200616/EW> MOST RECENT UPDATE WEEK: FILE COVERS 1978 TO DATE >>> IMAGES ARE AVAILABLE ONLINE AND FOR EMAIL-PRINTS <<< >>> NEW IPC8 DATA AND FUNCTIONALITY NOW AVAILABLE IN THIS FILE. http://www.stn-international.de/stndatabases/details/ipc-reform.html >>> >>> FOR CHANGES IN PCTFULL PLEASE SEE HELP CHANGE (last updated April 10, 2006) <<< => s losartan or enalapril 1362 LOSARTAN 1 LOSARTANS 1362 LOSARTAN (LOSARTAN OR LOSARTANS) 2445 ENALAPRIL 3 ENALAPRILS 2445 ENALAPRIL (ENALAPRIL OR ENALAPRILS) L18 2831 LOSARTAN OR ENALAPRIL

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22498 NEOPLAS?
L19
         96710 CANCER? OR TUMOR? OR NEOPLAS?
=> s 118 and 119
        1575 L18 AND L19
L20
=> s 118/clm
          284 LOSARTAN/CLM
           458 ENALAPRIL/CLM
L21
          571 (LOSARTAN/CLM OR ENALAPRIL/CLM)
=> s 119/clm
        22195 CANCER?/CLM
         15078 TUMOR?/CLM
         3646 NEOPLAS?/CLM
L22
        31673 (CANCER?/CLM OR TUMOR?/CLM OR NEOPLAS?/CLM)
=> s 121 and 122
        141 L21 AND L22
=> s 123 not py>2000
       592560 PY>2000
          15 L23 NOT PY>2000
L24
=> s 124 not py>1999
       672418 PY>1999
T<sub>2</sub>5
          12 L24 NOT PY>1999
=> d ibib 1-6
      ANSWER 1 OF 12
                       PCTFULL COPYRIGHT 2006 Univentio on STN
                       1999043663 PCTFULL ED 20020515
ACCESSION NUMBER:
TITLE (ENGLISH):
                       N-[(SUBSTITUTED FIVE-MEMBERED DI- OR TRIAZA
                       DIUNSATURATED RING) CARBONYL] GUANIDINE DERIVATIVES FOR
                       THE TREATMENT OF ISCHEMIA
TITLE (FRENCH):
                       DERIVES DE LA N-[(A CYCLE DI OU TRIAZA DIINSATURE
                       SUBSTITUE) CARBONYLE] GUANIDINE UTILISES POUR LE
                       TRAITEMENT DE L'ISCHEMIE
INVENTOR(S):
                       HAMANAKA, Ernest, S.;
                       GUZMAN-PEREZ, Angel;
                       RUGGERI, Roger, B.;
                       WESTER, Ronald, T.;
                       MULARSKI, Christian, J.
PATENT ASSIGNEE(S):
                       PFIZER PRODUCTS INC.;
                       HAMANAKA, Ernest, S.;
                       GUZMAN-PEREZ, Angel;
                       RUGGERI, Roger, B.;
                       WESTER, Ronald, T.;
                       MULARSKI, Christian, J.
LANGUAGE OF PUBL.:
                       English
DOCUMENT TYPE:
                       Patent
PATENT INFORMATION:
                       NUMBER
                                          KIND
                                                  DATE
                       ______
                       WO 9943663
                                            Al 19990902
DESIGNATED STATES
                       AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE
      W:
                       ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR
                       KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT
                       RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU
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ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT

64916 TUMOR?

SE BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG

APPLICATION INFO.: WO 1999-IB206 A 19990205 PRIORITY INFO.: US 1998-60/076,362 19980227

ANSWER 2 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1999030690 PCTFULL ED 20020515

ORAL DELIVERY FORMULATION TITLE (ENGLISH):

TITLE (FRENCH): FORMULATION D'ADMINISTRATION PAR VOIE ORALE

INVENTOR(S): COMPTON, Bruce, Jon; SOLARI, Nancy, E.; FLANAGAN, Margaret, A.

PATENT ASSIGNEE(S): AXIA THERAPEUTICS, INC.;

COMPTON, Bruce, Jon; SOLARI, Nancy, E.; FLANAGAN, Margaret, A.

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION:

NUMBER KIND DATE ______

WO 9930690 A1 19990624

DESIGNATED STATES W:

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT

SE BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG

WO 1998-US26627 A 19981215 US 1997-60/069,501 19971215 US 1998-60/073,867 19980204 US 1998-09/055,163 19980404 US 1998-09/055,560 19980406 APPLICATION INFO.: PRIORITY INFO.:

L25 ANSWER 3 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN

L25 ANSWER 3 OL 12

ACCESSION NUMBER: 1999018956 PCTFULL ED 2002

TITLE (ENGLISH): 12(S)-HETE RECEPTOR BLOCKERS

TITLE (FRENCH): INHIBITEURS DU RECEPTEUR DE 12(S)-HETE

MATARAJAN, Rama, Devi;

NADLER, Jerry, L.

PATENT ASSIGNEE(S): CITY OF HOPE
LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION: NUMBER KIND DATE

WO 9918956 A1 19990422

DESIGNATED STATES

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE W: ES FI GB GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE

CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ

CF CG CI CM GA GN GW ML MR NE SN TD TG

WO 1998-US21570 A 19981014 APPLICATION INFO.: PRIORITY INFO.: US 1997-60/062,335 19971015

ANSWER 4 OF 12 T₂5 PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1999008596 PCTFULL ED 20020515 TITLE (ENGLISH): MEASUREMENT OF CAPILLARY RELATED INTERSTITIAL FLUID USING ULTRASOUND METHODS AND DEVICES

TITLE (FRENCH): MESURE DU FLUIDE INTERSTITIEL PROPRE AUX CAPILLAIRES

UTILISANT DES METHODES ET DES DISPOSITIFS

ECHOGRAPHIQUES

INVENTOR(S): LANG, Philipp;

MENDLEIN, John, D.

LANG, Philipp; PATENT ASSIGNEE(S):

MENDLEIN, John, D.

LANGUAGE OF PUBL.: English DOCUMENT TYPE:

Patent

PATENT INFORMATION:

NUMBER KIND DATE _____

WO 9908596 A1 19990225

DESIGNATED STATES

W:

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF

BJ CF CG CI CM GA GN GW ML MR NE SN TD TG

WO 1998-US17238 A 19980819

APPLICATION INFO.: PRIORITY INFO.: US 1997-08/914,527 19970819

ANSWER 5 OF 12

PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1998051282 PCTFULL ED 20020514 .
TITLE (ENGLISH): SOLID POROUS MATRICES AND METHODS OF MAKING AND USING

THE SAME

TITLE (FRENCH): MATRICES POREUSES SOLIDES, LEUR PROCEDE DE FABRICATION

ET LEUR UTILISATION

L25

INVENTOR(S):

PATENT ASSIGNEE(S):

LANGUAGE OF PUBL.:

English

DOCUMENT TYPE:

DOCUMENT TYPE:

Patent

PATENT INFORMATION:

NUMBER KIND DATE _____

WO 9851282 A1 19981119

DESIGNATED STATES

W:

AU BR CA CN JP KR NZ AT BE CH CY DE DK ES FI FR GB GR

IE IT LU MC NL PT SE

APPLICATION INFO.: PRIORITY INFO.:

WO 1998-US9570 A 19980512

US 1997-60/046,379 19970513

US 1998-9/075,477

19980511

ANSWER 6 OF 12

PCTFULL COPYRIGHT 2006 Univentio on STN

TITLE (ENGLISH):

ACCESSION NUMBER: 1998036784 PCTFULL ED 20020514
TITLE (ENGLISH): COATED IMPLANTABLE MEDICAL DEVICE

TITLE (FRENCH):

DISPOSITIF MEDICAL IMPLANTABLE DOTE D'UN REVETEMENT

RAGHEB, Anthony, O.; INVENTOR(S): BATES, Brian, L.;

FEARNOT, Neal, E.; KOZMA, Thomas, G.;

VOORHEES, William, D., III;

GERSHLICK, Anthony, H.

PATENT ASSIGNEE(S):

COOK INCORPORATED

LANGUAGE OF PUBL.:

English

DOCUMENT TYPE:

Patent

PATENT INFORMATION:

NUMBER

KIND DATE ______

WO 9836784 A1 19980827

DESIGNATED STATES

W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE

ES FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE

CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF

CG CI CM GA GN ML MR NE SN TD TG

APPLICATION INFO.: WO 1998-US3438 A 19980220 PRIORITY INFO.: US 1997-60/038,459 19970220

=> d kwic 2

L25 ANSWER 2 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN

CLMEN. . . a drug and the drug is selected from

> the group consisting of- isotretinoin; oxazepain; lorazepam; piroxicam; loperamide;

bromopheniramine; phenylpropanolanime; loratadine; famotidine;

ordansetron; enalapril; captopril;

phloroglucinol; nicergoline; acetaminophen; metapimazine;

dihydroergotamine; fexofenadine-HCI

and albuterol.

The method of claim 24, wherein the subject is selected from the group consisting

of a geriatric subject, a subject with cancer, a subject who

is post-surgically recovering, a child and

a pregnant mother.

=> d ibib 7-12

ANSWER 7 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN L25

ACCESSION NUMBER: 1998032718 PCTFULL ED 20020514

TITLE (ENGLISH): NEW FATTY ACID DERIVATIVES

NOUVEAUX DERIVES D'ACIDE GRAS TITLE (FRENCH):

INVENTOR(S): MYHREN, Finn; BORRETZEN, Bernt;

DALEN, Are;

SANDVOLD, Marit, Liland

PATENT ASSIGNEE(S): NORSK HYDRO ASA; MYHREN, Finn;

BORRETZEN, Bernt;

DALEN, Are;

SANDVOLD, Marit, Liland

LANGUAGE OF PUBL.:

English Patent

DOCUMENT TYPE: PATENT INFORMATION:

NUMBER KTND DATE

WO 9832718 A1 19980730

DESIGNATED STATES

W:

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU

SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ

CF CG CI CM GA GN ML MR NE SN TD TG

A 19980123 APPLICATION INFO.: WO 1998-NO21 19970124 PRIORITY INFO.: GB 1997-9701441.9

ANSWER 8 OF 12 L25 PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1998032022 PCTFULL ED 20020514
TITLE (ENGLISH): GROWTH FACTOR-DEPENDENT DISEASES
TITLE (FRENCH): MALADIES LIEES AU FACTEUR DE CROISSANCE

TITLE (FRENCH):

INVENTOR(S): EPSTEIN, Richard, John

PATENT ASSIGNEE(S): IMPERIAL EXPLOITATION LIMITED;

EPSTEIN, Richard, John

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION:

NUMBER KIND DATE -----WO 9832022 A1 19980723

DESIGNATED STATES

JP US AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT W:

SE

APPLICATION INFO.: WO 1998-GB33 A 19980115

PRIORITY INFO.: GB 1997-9700933.6 19970117

ANSWER 9 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1998018610 PCTFULL ED 20020514

TITLE (ENGLISH): EMBEDDING AND ENCAPSULATION OF CONTROLLED RELEASE

PARTICLES

TITLE (FRENCH): INCLUSION ET ENCAPSULATION DE PARTICULES A LIBERATION

CONTROLEE

INVENTOR(S): VAN LENGERICH, Bernhard, H.

PATENT ASSIGNEE(S): VAN LENGERICH, Bernhard, H. LANGUAGE OF PUBL.: English

DOCUMENT TYPE: Patent

PATENT INFORMATION:

KIND DATE NUMBER _____ WO 9818610 A1 19980507

DESIGNATED STATES

AU CA JP NO PL US AT BE CH DE DK ES FI FR GB GR IE IT W: .

LU MC NL PT SE

LU MC NL PT SE WO 1997-US18984 A 19971027 US 1996-60/029,038 19961028 APPLICATION INFO.: PRIORITY INFO.: US 1997-60/052,717 19970716

L25 ANSWER 10 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN

ACCESSION NUMBER: 1998017331 PCTFULL ED 20020514 TITLE (ENGLISH): SILVER IMPLANTABLE MEDICAL DEVICE

TITLE (FRENCH): DISPOSITIF MEDICAL IMPLANTABLE ET CONTENANT DE L'ARGENT

INVENTOR(S): BATES, Brian, L.; OSBORNE, Thomas, A.; ROBERTS, Joseph, W.;

FEARNOT, Neal, E.; KOZMA, Thomas, G.; RAGHEB, Anthony, O.; VOORHEES, William, D., III

PATENT ASSIGNEE(S): COOK INCORPORATED; MED INSTITUTE, INC.

LANGUAGE OF PUBL.: English Patent

DOCUMENT TYPE: PATENT INFORMATION:

KIND NUMBER DATE WO 9817331 A1 19980430

DESIGNATED STATES

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE W: ES FI GB GE GH HU IL IS JP KE KG KP KR KZ LC LK LR LS

LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG

SI SK SL TJ TM TR TT UA UG UZ VN YU ZW GH KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA

GN ML MR NE SN TD TG

APPLICATION INFO.: WO 1997-US19188 A 19971023 US 1996-60/029,158 US 1996-8/741,565 US 1997-8/803,843 PRIORITY INFO.: 19961024 19961031 19970224

ANSWER 11 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN L25

ACCESSION NUMBER: 1997031654 PCTFULL ED 20020514

TITLE (ENGLISH): NITRIC OXIDE DONORS CAPABLE OF REDUCING TOXICITY FROM

DRUGS

TITLE (FRENCH): DONNEURS D'OXYDE NITRIQUE CAPABLES DE DIMINUER LA

TOXICITE DE MEDICAMENTS

DEL SOLDATO, Piero INVENTOR(S):

PATENT ASSIGNEE(S): NICOX S.A.;

DEL SOLDATO, Piero

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION:

NUMBER KIND DATE WO 9731654 Al 19970904

DESIGNATED STATES

W:

AL AU BB BG BR CA CN CZ EE GE HU IL IS JP KP KR LK LR LT LV MG MK MN MX NO NZ PL RO RU SG SI SK TR TT UA US UZ VN KE LS MW SD SZ UG AM AZ BY KG KZ MD RU TJ TM AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ

CF CG CI CM GA GN ML MR NE SN TD TG WO 1997-EP873 A 19970224

APPLICATION INFO.: PRIORITY INFO.: IT 1996-MI96A000352 19960226

ANSWER 12 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN T.25

ACCESSION NUMBER: 1990006775 PCTFULL ED 20020513
TITLE (ENGLISH): A NOVEL NONPHOSPHOLIPID LIPOSOME COMPOSITION FOR

SUSTAINED RELEASE OF DRUGS NOUVELLE COMPOSITION DE LIPOSOMES NON PHOSPHOLIPIDIQUE TITLE (FRENCH):

A LIBERATION SOUTENUE DE MEDICAMENTS

RADHAKRISHNAN, Ramachandran INVENTOR(S): PATENT ASSIGNEE(S): LIPOSOME TECHNOLOGY, INC.

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent .

PATENT INFORMATION:

NUMBER KIND DATE ______ WO 9006775 A1 19900628

DESIGNATED STATES

W:

AT AU BE CH DE DK ES FI FR GB IT JP LU NL NO SE

WO 1989-US5525 A 19891206 APPLICATION INFO .: US 1988-284,158 US 1988-284,216 19881214 PRIORITY INFO.: 19881214 US 1989-Not furnished 19891201

=> d kwic 12

ANSWER 12 OF 12 PCTFULL COPYRIGHT 2006 Univentio on STN L25

CLMEN. . . enviroxime,

ribavarin, rimantadine, amantadine, penicillin, erythromycin, tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycin, gentamycin, tobramycin,

piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B. micozanole, apresoline, atenolol, captopril, verapamil, enalapril, dopamine, dextroamphetamine, pentamidine, pyribenzamine, chlorpheniramine, diphenhydramine, interferon, interleukinenviroxime, ribavarin, rimantadine, amantadine, penicillin, erythromycin, tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycin, gentamycin, tobramycin, piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B, micozanole, apresoline, atenolol, captopril, verapamil, enalapril, dopamine, dextroamphetamine, pentamidine, pyribenzamine, chlorpheniramine, diphenhydramine, interferon, interleukinenviroxime, ribavarin, rimantadine, amantadine, penicillin, erythromycin, tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycin, gentamycin, tobramycin, piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B, micozanole, apresoline, atenolol, captopril, verapamil, enalapril, dopamine, dextroamphetamine, pentamidine, pyribenzamine, chlorpheniramine, diphenhydramine, interferon, interleukinenviroxime, ribavarin, rimantadine, amantadine, penicillin, erythromycin, F tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycint gentamycin, tobramycin, piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B, micozanole, apresoline, atenolol, captopril, verapamil, enalapril, dopaminer dextroamphetaxine, pentamidine, pyribenzamine, chlorpheniramine, diphenhydramine, interferon, interieukinenviroxime, ribavarin, rimantadine, amantadine, penicillin, erythromycinr tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycin, gentamycin, tobramycin, piperacillinr moxalactamr cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B, micozanole,, apresoline,, atenolol,, captopril, verapamil, enalapril, dopamine, dextroamphetamine, pentamidine, pyribenzamine, chlorpheniramine, diphenhydramine, interferon, interleukincholesterol sulfate, 40 mole % of cholesterol and 10 mole t of the drug of Claim 25. 28a A method of suppressing neoplastic growth by administering to a person in need of such treatment a therapeutically effective amount of nonconventional liposome composition comprising nonphospholipid lipids and. enviroxime, ribavarin, rimantadine, amantadine, penicillin, erythromycin, tetracyclin, cephalothin, cefotaxime, carbenicillin, vancomycin, gentamycin, tobramycin, piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin, amikacin, amphotericin B, micozanole, apresoline, atenolol, captopril, verapamil, enalapril, dopamine,

dextroamphetamine, pentamidine, pyribenzamine,

chlorpheniramine, diphenhydramine, interferon, interleukin-

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enviroxime,
       ribavarin, rimantadine, amantadine, penicillin,
       erythromycin, tetracyclin, cephalothin, cefotaxime,
       carbenicillin, vancomycin, gentamycin, tobramycin,
       piperacillin, moxalactam, cefazolin, cefadroxil, cefoxitin,
       amikacin, amphotericin B, micozanole, apresoline, atenolol,
       captopril, verapamil, enalapril, dopamine,
       dextroamphetamine, pentamidine, pyribenzamine,
       chlorpheniramine, diphenhydramine, interferon, interleukin-
=> d his
     (FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)
     FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006
               E "LOSARTAN"/CN 25
              1 S E3
                E "ENALAPRIL"/CN 25
              1 S E3
     FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006
           4684 S L1 OR L2
         704495 S CANCER? OR TUMOR? OR NEOPLAS?
            146 S L4 AND L3
             22 S L3 (L) L4
              3 S L6 NOT PY>2000
         266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?
           220 S L8 AND L3
             46 S L8 (L) L3
             18 S L10 NOT PY>2000
              7 S L10 AND L4
           7727 S LOSARTAN OR ENALAPRIL
             92 S L13 (L) L4
             28 S L14 NOT PY>2000
             23 S L14 NOT PY>1999
              4 S L16 AND L8
     FILE 'PCTFULL' ENTERED AT 10:25:28 ON 26 APR 2006
          2831 S LOSARTAN OR ENALAPRIL
          96710 S CANCER? OR TUMOR? OR NEOPLAS?
          1575 S L18 AND L19
            571 S L18/CLM
          31673 S L19/CLM
           141 S L21 AND L22
             15 S L23 NOT PY>2000
             12 S L24 NOT PY>1999
=> s 124 not 125
             3 L24 NOT L25
=> d ibib 1-3
      ANSWER 1 OF 3
                        PCTFULL
                                   COPYRIGHT 2006 Univentio on STN
                        2000074742 PCTFULL ED 20020515
ACCESSION NUMBER:
                        DEVICES AND COMPOUNDS FOR TREATING ARTERIAL RESTENOSIS
TITLE (ENGLISH):
TITLE (FRENCH):
                        DISPOSITIFS ET COMPOSES SERVANT A TRAITER LA RESTENOSE
```

ARTERIELLE INVENTOR(S): ZAHRADKA, PeterRP: ADE & COMPANY PATENT ASSIGNEE(S): CARDIO VASCULAR SOLUTIONS INC.;

ZAHRADKA, Peter LANGUAGE OF PUBL.: English

DOCUMENT TYPE: Patent

L1

L2

L3

L4L5

L6

L7

 \cdot L8 L9

L10

L11L12

L13

L14

L15 L16

L17

L18

L19

L20

L21 L22

L23

L24

L25

L26

PATENT INFORMATION:

NUMBER KIND DATE ------

WO 2000074742

A1 20001214

DESIGNATED STATES

W:

AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW GH GM KE LS MW MZ SD SL SZ TZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI

CM GA GN GW ML MR NE SN TD TG

APPLICATION INFO.:

WO 2000-CA653 A 20000601

US 1999-60/150,696 19990602

PRIORITY INFO.:

ANSWER 2 OF 3 L26 ACCESSION NUMBER: TITLE (ENGLISH):

PCTFULL COPYRIGHT 2006 Univentio on STN 2000040227 PCTFULL ED 20020515

METHODS FOR TREATING CONDITIONS ASSOCIATED WITH THE

TITLE (FRENCH):

ACCUMULATION OF EXCESS EXTRACELLULAR MATRIX

PROCEDES DE TRAITEMENT D'ETATS ASSOCIES A

L'ACCUMULATION D'UN EXCEDENT DE MATRICE EXTRACELLULAIRE

INVENTOR(S):

NOBLE, Nancy, A.; BORDER, Wayne, A.;

LAWRENCE, Daniel, A.

PATENT ASSIGNEE(S):

UNIVERSITY OF UTAH; AMERICAN NATIONAL RED CROSS;

NOBLE, Nancy, A.; BORDER, Wayne, A.; LAWRENCE, Daniel, A.

LANGUAGE OF PUBL.:

English

DOCUMENT TYPE:

Patent

PATENT INFORMATION:

KIND DATE NUMBER

WO 2000040227 A2 20000713

DESIGNATED STATES

W:

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW GH GM KE LS MW SD SL SZ TZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG

APPLICATION INFO.: PRIORITY INFO.:

WO 2000-US179 A 20000105 US 1999-60/114,795 19990105

L26 ANSWER 3 OF 3 PCTFULL COPYRIGHT 2006 Univentio on STN ACCESSION NUMBER: 2000001706 PCTFULL ED 20020515 TITLE (ENGLISH): N-TERMINAL SITE SELECTIVE INHIBITORS OF HUMAN

ANGIOTENSIN CONVERSION ENZYME (ACE)

TITLE (FRENCH): INVENTOR(S):

INHIBITEURS SELECTIFS DE SITE N-TERMINAL DE L'ECA

DIVE, Vincent; COTTON, Joel;

CUNIASSE, Philippe; YIOTAKIS, Athanasios;

CORVOL, Pierre; MICHAUD, Annie;

CHAUVET, Marie-Therese;

MENARD, Joel; EZAN, Eric

PATENT ASSIGNEE(S):

COMMISSARIAT A L'ENERGIE ATOMIQUE;

INSTITUT NATIONAL DE LA SANTE ET DE LA RECHERCHE

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DIVE, Vincent;
                       COTTON, Joel;
                       CUNIASSE, Philippe;
                       YIOTAKIS, Athanasios;
                       CORVOL, Pierre;
                       MICHAUD, Annie;
                       CHAUVET, Marie-Therese;
                       MENARD, Joel;
                       EZAN, Eric
LANGUAGE OF PUBL.:
                       French
DOCUMENT TYPE:
                       Patent
PATENT INFORMATION:
                       NUMBER
                                        KIND DATE
                       ------
                       WO 2000001706 A1 20000113
DESIGNATED STATES
                       CA JP US AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC
      W:
                       NL PT SE
APPLICATION INFO .:
                       WO 1999-FR1581
                                          A 19990701
PRIORITY INFO.:
                       FR 1998-98/08464
                                              19980702
=> d kwic 3
      ANSWER 3 OF 3 PCTFULL COPYRIGHT 2006 Univentio on STN
L26
CLMFR 10 Composition selon la revendication 8,
      dans laquelle le traitement est un traitement anti-
        cancereux.
      la revendication 11,
      dans laquelle le medicament est destine a reguler la
      proliferation des cellules souches hematopoietiques de
      patients soumis a un traitement anti-cancereux.
      CH3
      1 rooo)
      CAPTOPRIL HS-CH2-CH - C- N - CH- COOH
      il
      0
      соон снз
      1 1 fooe)
        ENALAPRIL CH2- CH2 c N - CH-COOH
      0
      FOSINOPRIL ]]](CH2)4 - p - CH2C CH- COOH
      Uri
      FI Ga 1
      % D'INHIBITION DU MUTANT N-TERMINAL ACTIF. . .
=> d his
     (FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)
    FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006
               E "LOSARTAN"/CN 25
             1 S E3
L1
               E "ENALAPRIL"/CN 25
             1 S E3
L2
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FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006

MEDICALE INSERM;

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L3
          4684 S L1 OR L2
L4
         704495 S CANCER? OR TUMOR? OR NEOPLAS?
L5
            146 S L4 AND L3
L6
             22 S L3 (L) L4
L7
             3 S L6 NOT PY>2000
L8
         266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?
           220 S L8 AND L3
L9
             46 S L8 (L) L3
L10
             18 S L10 NOT PY>2000
L11
L12
              7 S L10 AND L4
           7727 S LOSARTAN OR ENALAPRIL
L13
L14
             92 S L13 (L) L4
L15
             28 S L14 NOT PY>2000
L16
             23 S L14 NOT PY>1999
L17
              4 S L16 AND L8
     FILE 'PCTFULL' ENTERED AT 10:25:28 ON 26 APR 2006
          2831 S LOSARTAN OR ENALAPRIL
L18
          96710 S CANCER? OR TUMOR? OR NEOPLAS?
L19
L20
          1575 S L18 AND L19
L21
           571 S L18/CLM
L22
          31673 S L19/CLM
L23
           141 S L21 AND L22
L24
            15 S L23 NOT PY>2000
             12 S L24 NOT PY>1999
L25
L26
             3 S L24 NOT L25
=> s angiotensin II
          7881 ANGIOTENSIN
           128 ANGIOTENSINS
          7949 ANGIOTENSIN
                 (ANGIOTENSIN OR ANGIOTENSINS)
        348075 II
          6036 IIS
        349694 II
                 (II OR IIS)
L27
          2257 ANGIOTENSIN II
                (ANGIOTENSIN(W)II)
=> s 127/ab
           497 ANGIOTENSIN/AB
             1 ANGIOTENSINS/AB
           497 ANGIOTENSIN/AB
                 ((ANGIOTENSIN OR ANGIOTENSINS)/AB)
         22147 II/AB
            18 IIS/AB
         22161 II/AB
                 ((II OR IIS)/AB)
           295 (ANGIOTENSIN II/AB)
L28
                 ((ANGIOTENSIN(W)II)/AB)
=> s 128 and 119
        43 L28 AND L19
=> s 118 and 129
         26 L18 AND L29
L30
=> s 130 not py>2000
       592560 PY>2000
L31
            8 L30 NOT PY>2000
=> s 130 not py>1999
        672418 PY>1999
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=> d ibib 1-6

PCTFULL COPYRIGHT 2006 Univentio on STN ANSWER 1 OF 6 L32

ACCESSION NUMBER: 1998046266 PCTFULL ED 20020514
TITLE (ENGLISH): COMPOSITIONS CONTAINING AN ANGIO

COMPOSITIONS CONTAINING AN ANGIOTENSIN II ANTAGONIST TITLE (ENGLISH):

AND AN ANGIOTENSIN II AGONIST FOR USE IN THE TREATMENT

OF ERECTILE DYSFUNCTION

TITLE (FRENCH): COMPOSITIONS CONTENANT UN ANTAGONISTE DE L'ANGIOTENSINE

II ET UN AGONISTE DE L'ANGIOTENSINE II, DESTINEES AU

TRAITEMENT DES DYSERECTIONS

INVENTOR(S): KIFOR, Imre;

WILLIAMS, Gordon

PATENT ASSIGNEE(S): BRIGHAM & WOMEN'S HOSPITAL, INC.

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION:

KIND DATE NUMBER -----WO 9846266 A1 19981022

DESIGNATED STATES

CA JP AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT W:

APPLICATION INFO.: A 19980326 WO 1998-US5893

PRIORITY INFO.: US 1997-60/041,875 19970411

ANSWER 2 OF 6 PCTFULL COPYRIGHT 2006 Univentio on STN ACCESSION NUMBER:

1998046224 PCTFULL ED 20020514

TITLE (ENGLISH): COMPOSITIONS AND METHOD FOR TREATING BLADDER

DYSFUNCTION

TITLE (FRENCH): COMPOSITIONS ET PROCEDES POUR TRAITER LE

DYSFONCTIONNEMENT DE LA VESSIE

INVENTOR(S): KIFOR, Imre;

WILLIAMS, Gordon; SULLIVAN, Maryrose, P.

PATENT ASSIGNEE(S): BRIGHAM & WOMEN'S HOSPITAL, INC.

LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent

PATENT INFORMATION:

NUMBER KIND DATE ______ WO 9846224 A1 19981022

DESIGNATED STATES

INVENTOR(S):

CA JP AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT W:

SE

WO 1998-US5886 APPLICATION INFO.: A 19980326 PRIORITY INFO.: US 1997-60/041,874 19970411 US 1997-60/041,875 19970411

T.32 ANSWER 3 OF 6 PCTFULL COPYRIGHT 2006 Univentio on STN 1998018496 PCTFULL ED 20020514

ACCESSION NUMBER: TITLE (ENGLISH): CONTRAST AGENTS TITLE (FRENCH): AGENTS DE CONTRASTE

KLAVENESS, Jo; NAEVESTAD, Anne; CUTHBERTSON, Alan

PATENT ASSIGNEE(S): NYCOMED IMAGING AS;

COCKBAIN, Julian; KLAVENESS, Jo; NAEVESTAD, Anne; CUTHBERTSON, Alan

LANGUAGE OF PUBL.: English

DOCUMENT TYPE: Patent PATENT INFORMATION: NUMBER KIND DATE WO 9818496 A2 19980507 DESIGNATED STATES AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE W: ES FI GB GE GH HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW GH KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML MR NE SN TD TG WO 1997-GB2956 A 19971028 APPLICATION INFO.: GB 1996-9622368.0 19961028
GB 1996-9622365.6 19961028
GB 1996-9622364.9 19961028
GB 1996-9622369.8 19961028
GB 1996-9622366.4 19961028
GB 1996-9622367.2 19961028
GB 1997-9700699.3 19970115
GB 1997-9706063.6 19970324 PRIORITY INFO.: ANSWER 4 OF 6 PCTFULL COPYRIGHT 2006 Univentio on STN ACCESSION NUMBER: 1995029674 PCTFULL ED 20020514 TITLE (ENGLISH): A METHOD OF MODIFYING ANGIOTENSIN RECEPTOR ACTIVITY FOR TREATMENT OF PREMENSTRUAL SYNDROME AND MEDIATION OF PAIN TITLE (FRENCH): METHODE DE MODIFICATION DE L'ACTIVITE DU RECEPTEUR DE L'ANGIOTENSINE POUR LE TRAITEMENT DU SYNDROME PREMENSTRUEL ET DE LA DOULEUR INVENTOR(S): DePADOVA, Anthony, S. DePADOVA, Anthony, S. PATENT ASSIGNEE(S): LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent PATENT INFORMATION: NUMBER KIND DATE ______ WO 9529674 A1 19951109 DESIGNATED STATES AM AT AU BB BG BR BY CA CH CN CZ DE DK ES FI GB GE HU W: JP KE KG KP KR KZ LK LT LU LV MD MG MN MW NO NZ PL PT RO RU SD SE SI SK TJ TT UA US UZ VN KE MW SD SZ UG AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML MR NE SN TD TG APPLICATION INFO.: WO 1995-US5312 A 19950428 PRIORITY INFO.: US 1994-8/235,468 19940429 L32 ANSWER 5 OF 6 PCTFULL COPYRIGHT 2000 CM.

ACCESSION NUMBER: 1995006410 PCTFULL ED 20020514

ANTAGONIST OF ANGIOTENSIN II RECEPTORS

ANTAGONIST OF RECEPTEURS DE L'ANGIOTENSIN DE L'ANGIOTENSI PCTFULL COPYRIGHT 2006 Univentio on STN ANTAGONISTE DES RECEPTEURS DE L'ANGIOTENSINE II INVENTOR(S): GRISWOLD, Don, Edgar; WHARTON, John PATENT ASSIGNEE(S): SMITHKLINE BEECHAM CORPORATION; GRISWOLD, Don, Edgar; WHARTON, John DOCUMENT TYPE: Patent

NUMBER

WO 9506410

KIND DATE

A1 19950309

PATENT INFORMATION:

DESIGNATED STATES

W:

AU BB BG BR BY CA CN CZ FI HU JP KP KR KZ LK MG MN MW NO NZ PL RO RU SD SI SK UA US VN AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML

MR NE SN TD TG

APPLICATION INFO.: PRIORITY INFO.:

WO 1994-US10258 US 1993-8/115,968 A 19940901 19930901

ANSWER 6 OF 6 ACCESSION NUMBER:

COPYRIGHT 2006 Univentio on STN PCTFULL

1992020661 PCTFULL ED 20020513

TITLE (ENGLISH): TITLE (FRENCH): INVENTOR(S):

N, N-DIACYLPIPERAZINES N, N-DIACYLPIPERAZINES ASHTON, Wallace, T.; GREENLEE, William, J.;

WU, Mu, Tsu; DORN, Conrad, P.;

MacCOSS, Malcolm; MILLS, Sander, G. MERCK & CO., INC.

PATENT ASSIGNEE(S): LANGUAGE OF PUBL.: DOCUMENT TYPE:

PATENT INFORMATION:

English Patent

NUMBER KIND DATE WO 9220661 A1 19921126

DESIGNATED STATES

W:

AT BE CA CH DE DK ES FR GB GR IT JP LU MC NL SE

APPLICATION INFO.: WO 1992-US4189 A 19920519 US 1991-703,953 PRIORITY INFO.: 19910522 US 1992-885,416 19920519

=> d kwic 5

L32 ANSWER 5 OF 6 PCTFULL COPYRIGHT 2006 Univentio on STN ABEN The present invention relates to the use of an angiotensin II receptor antagonist in the manufacture of a medicament for the treatment of chronic inflammatory disease states.

DETD . . for

angiotensin in the regulation of tissue injury, proliferation and differen-tiation. As such this would include treatment of disorders such as tumor growth, i.e.

neoplastic transformation and growth/metasis, bone marrow maturation and differen-tiation, skin maturation and differentiation, and hepatocyte maturation and differentiation. Chronic inflammatory diseases would.

Vol. 183, pp 989-995 (1992), and is a member of the superfamily of G protein-coupled transmembrane receptors. AT1 antagonists such as losartan (Whitebread et al., supra) al., J. Med. Chem., Vol. 34, pp 1514-1517 (1991), sensitivity to reducing agents such as dithiothreitol (DTT) (Whitebread et.

published January 20, 1988. Preferred compounds included within this class of All receptor antagonists are (hydroxymethyl)-imidazole (also referred to as Losartan herein); or a pharmaceutically acceptable salt thereof

µ M PD123319 (AT1 receptors remain unblocked) or with 10 µ M µ M PD123319 (AT1 receptors remain unblocked) or with 10 µ M

losartan (AT2 receptors remain unblocked). Guanine nucleotide sensitivity of AII binding was assessed by coincubating sections with 0.25 nM [1251]AII and 1. Losartan was kindly provided by DuPont Merck, Wilmington, SmithKline Beecham, King of Prussia, PA, U.S.A., and PD123319 by Parke-Davis, Ann Arbor, MI,. with osteoarthritis and those with rheumatoid arthritis, specific ([1251](Sar1, Ile8) AII binding to each structure was completely inhibited by the AT1 antagonist losartan (10 µM), but was not significantly inhibited by the specific AT2 antagonist PD123319 (10 µ M) (table 6) . Specific binding to. 163, pp 264- 291 (1989); Weinstock et al., J. Med. Chem., Vol. 34, 1514-1517 (1991). Binding studies performed in the presence of losartan or PD123319 provide no evidence for the presence of AT2 receptors in synovium from any of these disease groups. The data. . . enzyme inhibitors are listed. No patient had received glucocorticosteroids within the month prior to surgery. Abbreviations; AZA; azathioprine, CC; chondrocalcinosis, ENA; enalapril, MTX; methotrexate, OA; osteoarthritis, RA; rheumatoid arthritis, SZ; sulphasalazine 2.00) c 3.39 (2.19 to 5.25) Αl d 12.6 (6.6 to 24.6) d 8.32 (4.17 to 16.2) d Αl 15.1 (7.41 to 31.6) e 37.2 (15.9 to 87.1) e 102.3 (33.1 to Losartan 316.2)e 75.9 (32.4 to 177.8 1-(4carboxyphenyl)methyl)-1 H-imidazol-5-y1]-2(2-thienyl)-PD 123319. . . 10,000 nHill val= (Sarl, lle8) 1.0(0.8 to 1.3) 1.2(0.9 to 1.6) 1.6(1.2 to2.0) All All 1.0(0.6 to 1.5) 0.9(0.5 to 1.2)0.8(0.5 to)1.2) Losartan 0.6(0.4 to 0.8)0.9(0.5 to 1.2)0.6(0.3 to 0.8)1-(4carboxyphenyl)methyl)-1H-(2-thienyl)methyl-2-propenoic acid PD 123319 Footnotes to Table 5: a. . . c (Sarl, Ile8) All was significantly more potent than All (each p lt; 0.05), d All

Effect of the specific ATl antagonist Losartan and AT2 antagonist PD 123319 on binding of [1251] (Sarl, Ile8) angiotensin II bindin to human s novium

significant differences were observed in Ki.

was significantly more potent than losartan in lining cells 2-propenoic acid in all structures (each p lt; 0.05). e no

Total Nonspecific Losartan PD123319 Blood Vessels ОА 7.9 (6.0 to 10) 2.4 (2.2 to 2.6) a 2.9 (2.4 to 3.5) 3.0 (2.3 to 4.0) 1.6 (1.5 to 1.7) a 1.5 (1.5 to 1.6) b 2.9 (2.1 to 4.1 a Losartan (10 µ M) significantly inhibits binding to structure in synovium from patients with osteoarthritis (OA) or rheumatoid arthritis (RA) (each. . . => d ibib kwic 5-6 PCTFULL COPYRIGHT 2006 Univentio on STN ANSWER 5 OF 6 1995006410 PCTFULL ED 20020514 ACCESSION NUMBER: TITLE (ENGLISH): ANTAGONIST OF ANGIOTENSIN II RECEPTORS TITLE (FRENCH): ANTAGONISTE DES RECEPTEURS DE L'ANGIOTENSINE II INVENTOR(S): GRISWOLD, Don, Edgar; WHARTON, John PATENT ASSIGNEE(S): SMITHKLINE BEECHAM CORPORATION; GRISWOLD, Don, Edgar; WHARTON, John DOCUMENT TYPE: Patent PATENT INFORMATION: KIND DATE NUMBER _____ WO 9506410 A1 19950309 DESIGNATED STATES W: AU BB BG BR BY CA CN CZ FI HU JP KP KR KZ LK MG MN MW NO NZ PL RO RU SD SI SK UA US VN AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML MR NE SN TD TG A 19940901 APPLICATION INFO .: WO 1994-US10258 PRIORITY INFO.: US 1993-8/115,968 19930901 The present invention relates to the use of an angiotensin II receptor antagonist in the manufacture of a medicament for the treatment of chronic inflammatory disease states. DETD . . for angiotensin in the regulation of tissue injury, proliferation and differen-tiation. As such this would include treatment of disorders such as tumor growth, i.e. neoplastic transformation and growth/metasis, bone marrow maturation and differen-tiation, skin maturation and differentiation, and hepatocyte maturation and differentiation. Chronic inflammatory diseases would. . Vol. 183, pp 989-995 (1992), and is a member of the superfamily of G protein-coupled transmembrane receptors. AT1 antagonists such as losartan (Whitebread et al., supra) al., J. Med. Chem., Vol. 34, pp 1514-1517 (1991), sensitivity to reducing agents such as dithiothreitol (DTT) (Whitebread et. published January 20, 1988. Preferred compounds included within this class of All receptor antagonists are (hydroxymethyl)-imidazole (also referred to as Losartan herein); or a pharmaceutically acceptable salt thereof

µ M PD123319 (AT1 receptors remain unblocked) or with 10 µ M µ M PD123319 (AT1 receptors remain unblocked) or with 10 µ M

T.32

ABEN

losartan (AT2 receptors remain unblocked). Guanine nucleotide sensitivity of AII binding was assessed by coincubating sections with 0.25 nM [1251]AII and 1. Losartan was kindly provided by DuPont Merck, Wilmington, SmithKline Beecham, King of Prussia, PA, U.S.A., and PD123319 by Parke-Davis, Ann Arbor, MI,. with osteoarthritis and those with rheumatoid arthritis, specific ([1251](Sar1, Ile8)AII binding to each structure was completely inhibited by the AT1 antagonist losartan (10 µ M), but was not significantly inhibited by the specific AT2 antagonist PD123319 (10 µ M) (table 6) . Specific binding to. 163, pp 264- 291 (1989); Weinstock et al., J. Med. Chem., Vol. 34, 1514-1517 (1991). Binding studies performed in the presence of losartan or PD123319 provide no evidence for the presence of AT2 receptors in synovium from any of these disease groups. The data. . . enzyme inhibitors are listed. No patient had received glucocorticosteroids within the month prior to surgery. Abbreviations; AZA; azathioprine, CC; chondrocalcinosis, ENA; enalapril, MTX; methotrexate, OA; osteoarthritis, RA; rheumatoid arthritis, SZ; sulphasalazine 2.00) c 3.39 (2.19 to 5.25) Αl d 12.6 (6.6 to 24.6) d 8.32 (4.17 to 16.2) d Al 15.1 (7.41 to 31.6) e 37.2 (15.9 to 87.1) e 102.3 (33.1 to Losartan 316.2)e 75.9 (32.4 to 177.8 1-(4carboxyphenyl)methyl)-1 H-imidazol-5-y1]-2(2-thienyl)-PD 123319. . 10,000 nHill val= (Sarl, lle8) 1.0(0.8 to 1.3) 1.2(0.9 to 1.6) 1.6(1.2 to2.0) All All 1.0(0.6 to 1.5) 0.9(0.5 to 1.2)0.8(0.5 to)1.2) Losartan 0.6(0.4 to 0.8)0.9(0.5 to 1.2)0.6(0.3 to 0.8)1-(4carboxyphenyl)methyl)-1H-(2-thienyl) methyl-2-propenoic acid PD 123319 Footnotes to Table 5: . . c (Sar1, Ile8) All was significantly more potent than All (each p lt; 0.05), d All was significantly more potent than losartan in lining cells

Effect of the specific AT1 antagonist Losartan and AT2 antagonist PD 123319 on binding of [1251] (Sar1, Ile8) angiotensin II bindin to human s novium

significant differences were observed in Ki. 1.

2-propenoic acid in all structures (each p lt; 0.05), e no

```
Total
                            Nonspecific
                                                 Losartan
      PD123319
      Blood Vessels
                              ОА
                                      7.9 (6.0 to 10)
                                                            2.4 (2.2 to 2.6)
      a 2.9 (2.4 to 3.5)
                            b 7.1 (5.1. . . 2.6
               3.0 (2.3 to 4.0)
                                     1.6 (1.5 to 1.7)
                                                           a 1.5 (1.5 to 1.6)
      b 2.9 (2.1 to 4.1
      a Losartan (10 µ M) significantly inhibits binding to
      structure in synovium from patients with osteoarthritis (OA)
      or rheumatoid arthritis (RA) (each. .
      ANSWER 6 OF 6
                       PCTFULL COPYRIGHT 2006 Univentio on STN
ACCESSION NUMBER:
                       1992020661 PCTFULL ED 20020513
TITLE (ENGLISH):
                       N, N-DIACYLPIPERAZINES
                       N, N-DIACYLPIPERAZINES
TITLE (FRENCH):
INVENTOR(S):
                       ASHTON, Wallace, T.;
                       GREENLEE, William, J.;
                       WU, Mu, Tsu;
                       DORN, Conrad, P.;
                       MacCOSS, Malcolm;
                       MILLS, Sander, G.
PATENT ASSIGNEE(S):
                       MERCK & CO., INC.
LANGUAGE OF PUBL.:
                       English
DOCUMENT TYPE:
                       Patent
PATENT INFORMATION:
                       NUMBER KIND DATE
                       WO 9220661 A1 19921126
DESIGNATED STATES
      W:
                       AT BE CA CH DE DK ES FR GB GR IT JP LU MC NL SE
                       WO 1992-US4189 A 19920519
APPLICATION INFO.:
                       US 1991-703,953
PRIORITY INFO.:
                                               19910522
                       US 1991-703,953 19910522
US 1992-885,416 19920519
      Diacylpiperazines of general structure (I) are: angiotensin
ABEN
      II (A-II) antagonists selective for
      the type 2 (AT2) subtype useful in the treatment of cerebrovascular,
       cognitive, and CNS disorders;
      tachykinin. .
DETD
      . . . processes leading to ovulation has
      been reviewed (Andrade-Gordon, et al. Biochem,
      Pharmacol., 42, 715-719 (1991)),
      1 5
      In addition, AT2 receptors are found in
      neuronal tumor cells (Speth, et al, Peptide Res,, 2,
      232-239 (1989)) and in transformed human neural cells
      (Tallant, et al, Llypertension, 17, 1135-1143 (1991)).
    of angiotensin II to
      AT2 receptors in reproductive organs. The compounds
      of the present invention are also useful as
      anticancer agents for brain cancers and other cancers
      wherein the AT2 receptor is prevelant,
      SUBSTANCE P ANTAGONISM ASSAY
      The compounds of this invention are useful
      for antagonizing substance P in the treatment. .
      the hypertensive eye in solution in a suitable
      ophthalmic vehicle, Also, these compounds may be
      useful in the reversal of multidrug resistance in
        tumor cells by enhancing the efficacy -of
       chemotherapeutic agents. In add ition, these
```

compounds may have activity in blocking calcium channels in insect brain. . .

These include acetylcholinesterase inhibitors such as heptylphysostigmine and tetrahydroacridine (THA; tacrine), muscarinic agonists such as oxotremorine, inhibitors of angiotensin-converting enzyme such as octylramipril, captopril, ceranapril, enalapril, lisinopril, fosinopril and zofenopril, centrally-acting calcium channel blockers and as nimodipine, and nootropic agents such as piracetam.

CLMEN 22 A method for inhibiting the growth of neruonal tumor cells which contain AT2 receptors in a patient in need thereof which comprises the administration to the patient of an effective amount of. . .

35 A method for the treatment of multidrug resistant tumor cells comprising the administration to a mammal in need of such treatment of an effective amount of the compound of Claim 1,

=> d his

(FILE 'HOME' ENTERED AT 10:11:24 ON 26 APR 2006)

FILE 'REGISTRY' ENTERED AT 10:11:33 ON 26 APR 2006 E "LOSARTAN"/CN 25

L1 1 S E3

E "ENALAPRIL"/CN 25

L2 1 S E3

FILE 'CAPLUS' ENTERED AT 10:12:56 ON 26 APR 2006

L3 4684 S L1 OR L2

L4 704495 S CANCER? OR TUMOR? OR NEOPLAS?

L5 146 S L4 AND L3

L6 22 S L3 (L) L4

L7 3 S L6 NOT PY>2000

L8 266785 S PROLIF? OR ANGIOGEN? OR NEOVASCULAR?

L9 220 S L8 AND L3

L10 46 S L8 (L) L3

L11 18 S L10 NOT PY>2000

L12 7 S L10 AND L4

L13 7727 S LOSARTAN OR ENALAPRIL

L14 92 S L13 (L) L4

L15 28 S L14 NOT PY>2000

L16 23 S L14 NOT PY>1999

L17 4 S L16 AND L8

FILE 'PCTFULL' ENTERED AT 10:25:28 ON 26 APR 2006

2831 S LOSARTAN OR ENALAPRIL

L19 96710 S CANCER? OR TUMOR? OR NEOPLAS?

L20 1575 S L18 AND L19

L21 571 S L18/CLM

L18

L24

L22 31673 S L19/CLM

L23 141 S L21 AND L22

15 S L23 NOT PY>2000

L25 12 S L24 NOT PY>1999

L26 3 S L24 NOT L25

L27 2257 S ANGIOTENSIN II

L28 295 S L27/AB

L29	43	S	L28	AND	L19
L30	26	S	L18	AND	L29
L31	8	S	L30	NOT	PY>2000
L32	6	s	L30	NOT	PY>1999